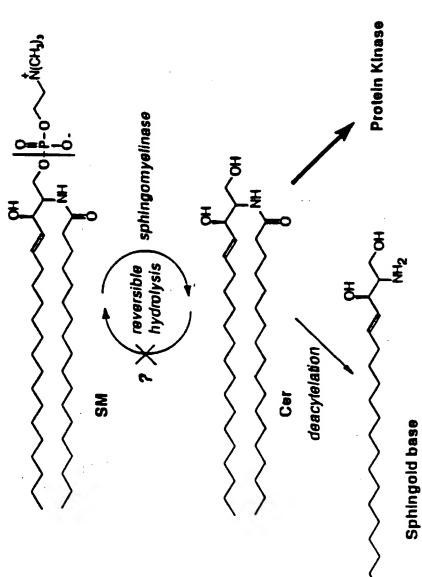


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(inhibitors of protein kinase C)



Fig. 2a

Type III Cer-1-TBDMS

C6 Cer-1-TBDMS

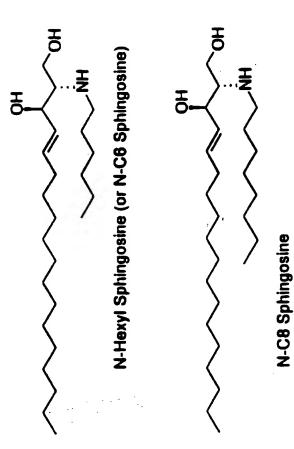
Fig. 2b

COTTOR LACTION

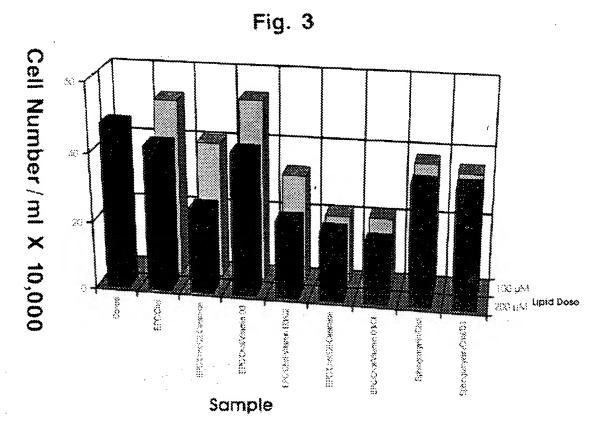
4,5-Diol C6-Cer

eszer tesete



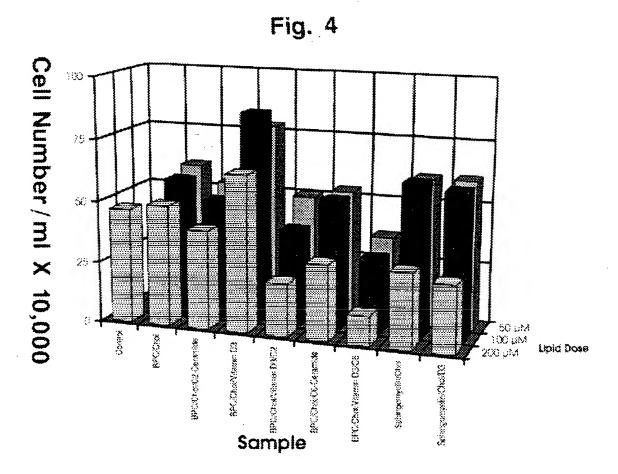


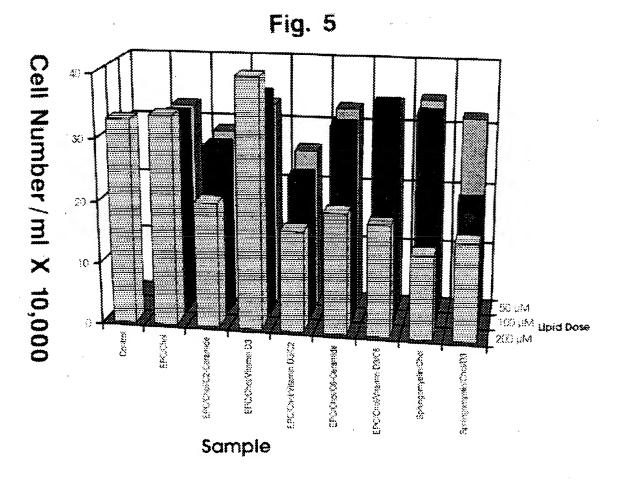
N-C10 Sphingoshne

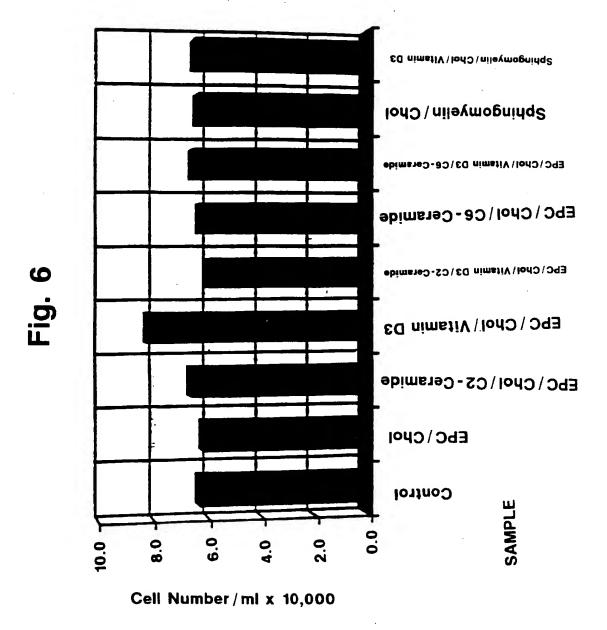


5,681,589



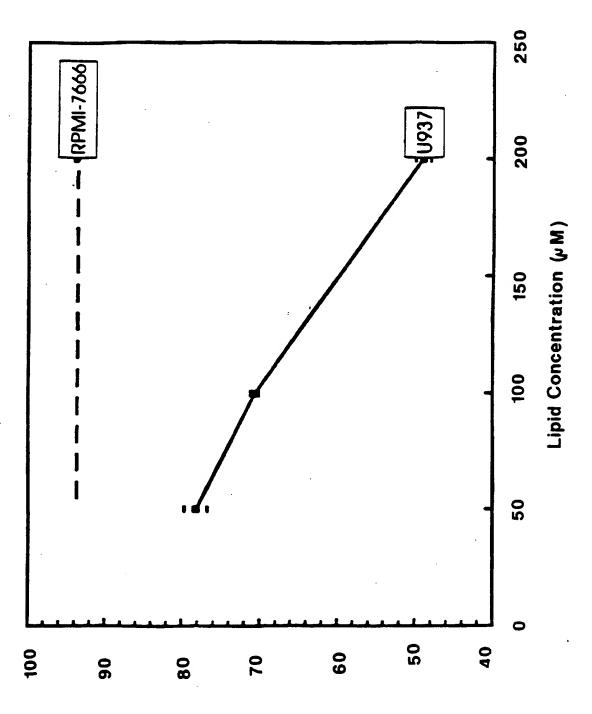






COTPOSOL LORYSO

Fig. 7

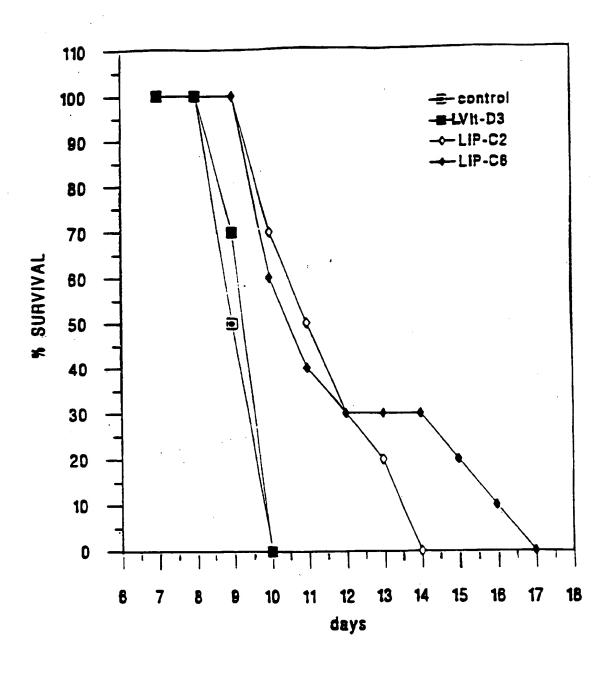


Cell Number as % Control



Sheet 11 of 21

Fig. 8



OGKESSY. 102799

In Vitro Sensitivity of A549 to C6-Ceramide Derivatives After Fig. 9A

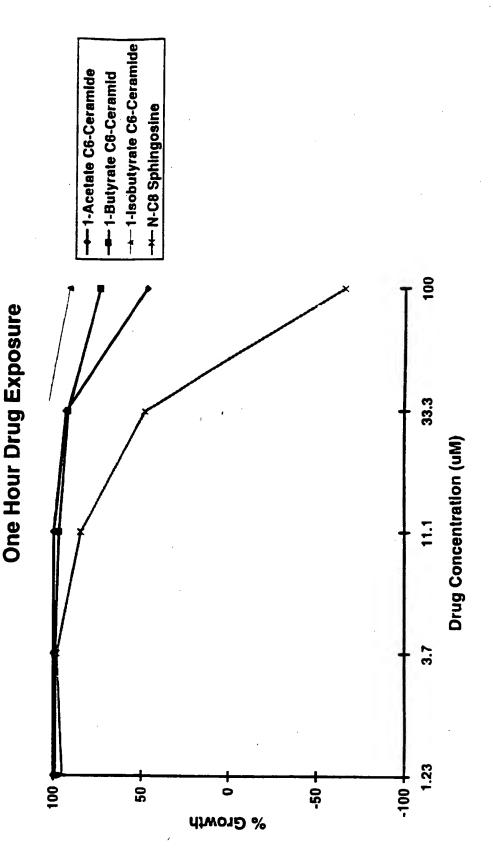


Fig. 9B

-4-1-Isobutyrate C6-C ramide -1-Butyrate C6-Ceramide -+- 1-Acetate C6-Ceramide --- N-C8 Sphingosine In Vitro Sensitivity of A549 to C6-Ceramide Derivatives After 100 Four Hour Drug Exposure 33.3 Drug Concentration (uM) 3.7 1.23 20 % Growth

DOYNOR LOUVE

Fig. 9C

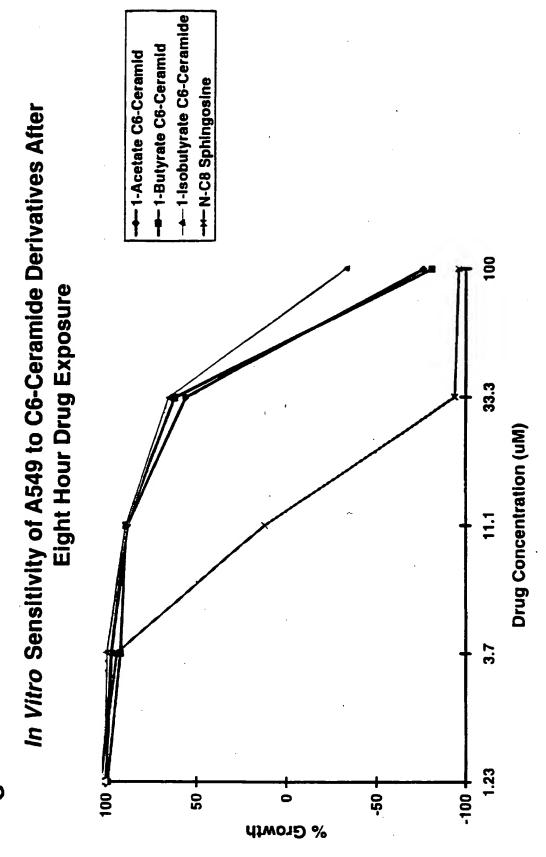


Fig. 9D

In Vitro Sensitivity of A549 to C6-Ceramide Derivatives After 24 Hour Drug Exposure

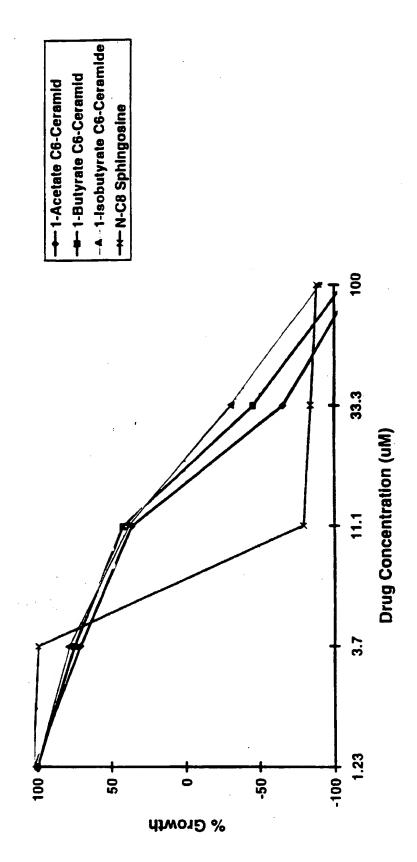


Fig. 9E

Doradat Tuese

In Vitro Sensitivity of A549 to C6-Ceramide Derivatives After 48 Hour Drug Exposure

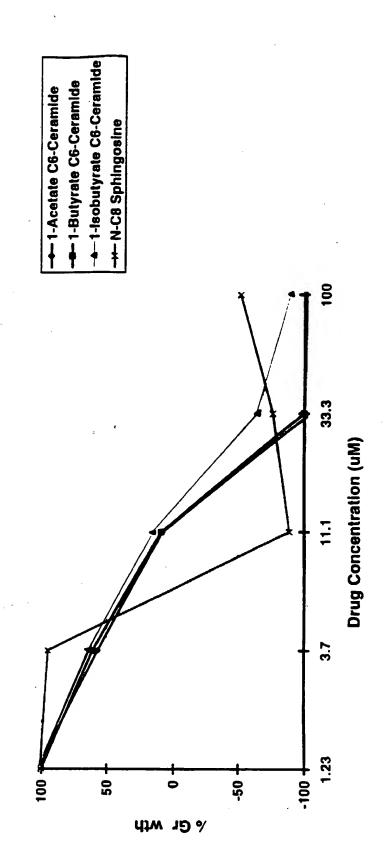
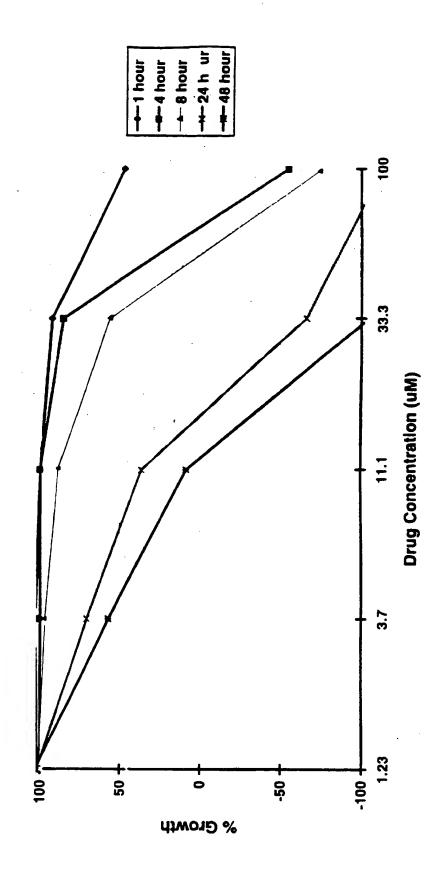


Fig. 10

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In Vitro Sensitivity of A549 to 1-Acetate C6-Ceramide at Different Exposure Times



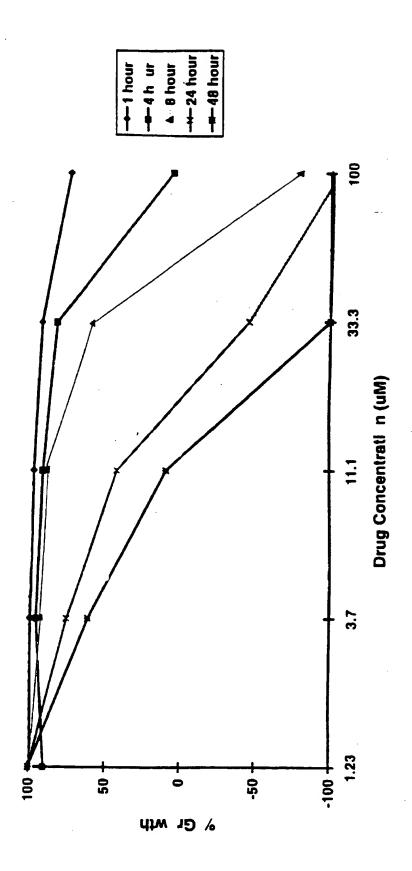
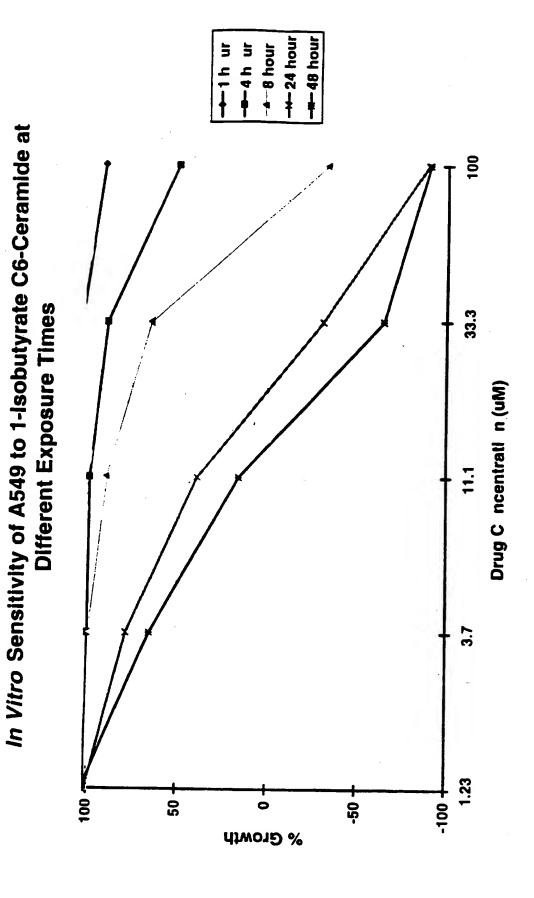


Fig. 1

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Fig. 12

COTUCUL TOURS



In Vitro Sensitivity of A549 to N-C8 Sphingosine at Different **Exposure Times** Fig. 13

